

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property  
Organization  
International Bureau



(43) International Publication Date  
21 October 2004 (21.10.2004)

PCT

(10) International Publication Number  
**WO 2004/090636 A1**

(51) International Patent Classification<sup>7</sup>: G03F 7/027

(21) International Application Number:  
PCT/KR2004/000860

(22) International Filing Date: 14 April 2004 (14.04.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
10-2003-0023432 14 April 2003 (14.04.2003) KR

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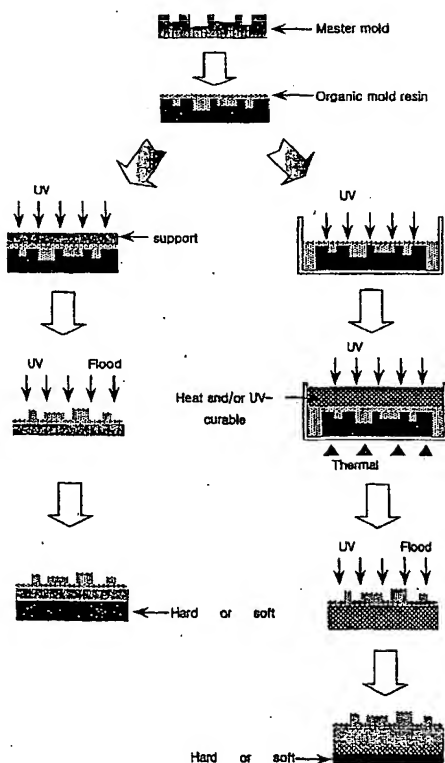
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(81) Designated States (unless otherwise indicated, for every  
kind of national protection available): AE, AG, AL, AM,  
AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,  
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,  
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,  
KG, KP, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG,  
MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH,  
PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,  
TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every  
kind of regional protection available): ARIPO (BW, GH,  
GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),

[Continued on next page]

(54) Title: RESIN COMPOSITION FOR MOLD USED IN FORMING MICROPATTERN, AND METHOD FOR FABRICATING  
ORGANIC MOLD THEREFROM



(57) Abstract: A resin composition for a mold used in forming micropatterns comprises (A) 40 to 90 parts by weight of an active energy curable urethane-based oligomer having a reactive group; (B) 10 to 60 parts by weight of a monomer reactive with the urethane-based oligomer; (C) 0.01 to 200 parts by weight of a silicone or fluorine containing compound, based on 100 parts of the sum of the components (A) and (B); and (D) 0.1 to 10 parts by weight of a photoinitiator, based on 100 parts of the sum of the components (A), (B) and (C). The inventive resin composition can be easily cured by the action of an active energy ray, and the organic mold fabricated therefrom is easily lifted off from a master without irreversible adhesion or generation of defects and have excellent dimensional and chemical stabilities.

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*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

**Published:**

— with international search report

**RESIN COMPOSITION FOR MOLD USED IN FORMING  
MICROPATTERN, AND METHOD FOR FABRICATING ORGANIC  
MOLD THEREFROM**

5

**FIELD OF THE INVENTION**

The present invention relates to a resin composition for a mold used in forming micropatterns, and to a method for fabricating an organic mold using same and the organic mold fabricated thereby.

10

**BACKGROUND OF THE INVENTION**

Various devices including integral circuits, semiconductors, electronic, photoelectric, display, magnetic or electromechanical devices and optical lens (e.g., prism sheet and lenticular lens sheet) involve micropatterns which have been conventionally formed by photolithography. However, the photolithography requires a complicated patterning process and a high production cost, and is not suitable for forming ultramicropatterns having a line width of below 100 nm.

Therefore, there has been recently developed a nano-imprint lithography wherein a pattern of a hard mold, e.g., a Si mold, is replicated on a thermoplastic polymer layer. This method is advantageous in fabricating a pattern having a narrow line width of about 7 nm owing to the hardness of the mold (see S.Y. Chou et al., J. Vac. Sci. Technol. B15, 2897(1997)). However, this nano-imprint lithography has the problems that the mold is not easily lifted off from the substrate and it may break during the pressurizing procedure under a high temperature and pressure condition.

Other non-traditional lithographic methods for fabricating micropatterns include micro-contact printing (mCP), micro-molding in capillaries (MIMIC), micro-transfer molding (mTM), soft molding, and capillary force lithography (CFL) methods. These methods generally employ a mold made of an elastic

polymer such as polydimethylsiloxane (PDMS), which is a silicon rubber type, but the PDMS mold has poor dimensional and chemical stabilities so that it cannot achieve a narrow pattern width of less than 500 nm.

Modified PDMS's including h-PDMS (hard PDMS) and hv-PDMS (photo-curable PDMS) have been developed to solve the problem of PDMS. However, h-PDMS still suffers from brittleness, low elongation at break, and poor conformal contact with a substrate (see Odom, Y. W. et al, *Langmuir*, 18, 5314-5320 (2002)), and hv-PDMS does not have enough modulus to replicate fine pattern below 100 nm even though produced to overcome the aforementioned some limitations of conventional PDMS (see *J. Am. Chem. Soc.* 125, 4060-4061 (2003)).

### SUMMARY OF THE INVENTION

Accordingly, it is an object of the present invention to provide a novel mold material having a modulus high enough for forming submicro-patterns and good chemical and dimensional stabilities which can be used for producing an organic mold that can be easily and repeatedly lifted off from a substrate without irreversible adhesion or generation of defects.

Further, it is another object of the present invention to provide a method for fabricating a mold using said material, and the mold fabricated thereby.

In accordance with one aspect of the present invention, there is provided a resin composition for a mold used in forming micropatterns, which comprises:

- (A) 40 to 90 parts by weight of an active energy curable urethane-based oligomer having a reactive group selected from the group consisting of (meth)acrylate, vinyl ether, arylether, and a combination thereof;
- (B) 10 to 60 parts by weight of a monomer (i.e., a reactive diluent) which is reactive with the urethane-based oligomer, and which has a reactive group selected from the group consisting of (meth)acrylate, vinyl ether, arylether, and a combination thereof;

(C) 0.01 to 200 parts by weight of a silicone or fluorine-containing compound (i.e., a functionalized additive), based on 100 parts of the sum of the components (A) and (B); and

5 (D) 0.1 to 10 parts by weight of a photo-initiator, based on 100 parts of the sum of the components (A), (B) and (C).

In accordance with another aspect of the present invention, there is provided a method for fabricating an organic mold, which comprises coating or casting the inventive resin composition on a pattern face of a mastermold, placing a support on the resin layer, irradiating an active energy ray to the resulting  
10 laminate to preliminarily cure the resin layer, lifting off the organic mold having a reverse pattern face to that of the mastermold and integrally formed with the support from the mastermold, and completely curing the organic mold.

In accordance with a further aspect of the present invention, there is provided a method for fabricating an organic mold, which comprises coating or  
15 casting the inventive resin composition on a pattern face of a mastermold, irradiating the resin layer with an active energy ray to preliminarily cure it, pouring a UV- or heat-curable resin composition onto the cured resin layer as a backbone, heating or irradiating the resultant to completely cure the resin layer and the backbone layer, lifting off the organic mold having a reverse pattern face  
20 to that of the mastermold and integrally formed with the backbone layer from the mastermold, and completely curing the organic mold.

In accordance with a still further aspect of the present invention, there is provided an organic mold having submicro-pattens, fabricated by any one of the inventive methods.

25

### **BRIEF DESCRIPTION OF THE DRAWINGS**

The above and other objects and features of the present invention will become apparent from the following description of the invention, when taken in  
30 conjunction with the accompanying drawings, which respectively show:

FIG. 1 : a schematic view showing the procedure for preparing a mold using a resin composition according to the present invention;

FIG. 2a : an electronic microscopic photograph of a mold having an equal line width and space of 80 nm fabricated by the inventive method;

5        FIG. 2b : an electronic microscopic photograph of a pattern formed on a substrate using the mold of FIG. 2a;

FIG. 3a : an electronic microscopic photograph of a mold having a cylindrical pattern fabricated by the inventive method;

10       FIG. 3b : an electronic microscopic photograph of a pattern formed on a substrate using the mold of FIG. 3a;

FIG. 4a : an optical microscopic photograph of a mastermold having a pyramid shaped pattern for use in fabricating a mold according to Comparative Example;

15       FIG. 4b : an optical microscopic photograph of an organic mold fabricated using the mastermold of FIG. 4a.

### **DETAILED DESCRIPTION OF THE INVENTION**

20       The present invention provides a resin composition for an organic mold to be used in forming micro-patterns, preferably submicro-patterns, and a method for fabricating the organic mold using said composition and the mold fabricated thereby.

25       The inventive resin composition is characterized by comprising an energy-curable components simultaneously having a hard segment such as cycloaliphatic or aromatic ring and a soft segment such as a linear aliphatic long chain, which makes it possible to form micropatterns and to impart the flexibility to a mold, and a silicon or fluorine containing compound which can provide a mold with an excellent releasing property while maintaining good physical properties of the mold. The compactness, i.e., the degree of cross-  
30       linkage in the molecular structure of the mold, may be enhanced by adjusting

the total amount of the reactive groups present in the composition.

Herein, the term “(meth)acrylate” is meant to represent acrylate and methacrylate, and the term “active energy ray” is meant to be an ultraviolet ray, infrared ray or electronic beam.

5 Specifically, the inventive resin composition comprises an active energy ray-curable urethane-based oligomer (“Component A”) for imparting a high elasticity and bendability to the composition. The active energy curable urethane-based oligomer may be a linear aliphatic, cycloaliphatic or aromatic urethane-based oligomer having at least two reactive groups, and a mixture  
10 thereof.

The urethane-based oligomer may be used in an amount of 40 to 90 parts by weight in the inventive composition. When the amount is less than the lower limit, the mechanical strength of a mold obtained from the composition becomes inferior, while when the amount exceeds the upper limit,  
15 the mold becomes too brittle.

The urethane-based oligomer may be partially substituted by a reactive oligomer for enhancing the physical properties such as flexibility, surface hardness, abrasive resistance, thermal resistance, weather resistance and chemical resistance of the composition. Such an optional oligomer may be a  
20 (meth)acrylated polyester, (meth)acrylated polyether, (meth)acrylated epoxy, (meth)acrylated polycarbonate, (meth)acrylated polybutadiene, or a mixture thereof, and may be employed in an amount of 0 to 200 % by weight of the urethane-based oligomer.

In the present invention, a monomer having the reactivity with the  
25 urethane-based oligomer (“Component B”) is employed as a reactive diluent, and representative examples thereof include (meth)acrylates such as isobornyl acrylate, 1,6-hexanediol acrylate, triethyleneglycol di(meth)acrylate, trimethylol propane triacrylate, tetraethyleneglycol di(meth)acrylate, 1,3-butanediol diacrylate, 1,4-butanediol diacrylate, diethyleneglycol diacrylate,  
30 neopentylglycol diacrylate, neopentylglycol di(meth)acrylate,

polyethyleneglycol di(meth)acrylate, pentaerythritol triacrylate, dipentaerythritol (hydroxy) pentaacrylate, alkoxyated tetraacrylate, octadecyl acrylate, isodecyl acrylate, lauryl acrylate, stearyl acrylate, behenyl acrylate, styrenic monomer, and a mixture thereof; and vinyl ethers and aryl ethers such as cyclohexyl vinyl ether, 2-ethylhexyl vinyl ether, dodecyl vinyl ether, aryl propyl ether, aryl butyl ether, 1,4-butanediol divinyl ether, 1,4-hexanediol divinyl ether, diethylene glycol divinyl ether, ethyleneglycol butyl vinyl ether, ethyleneglycol divinyl ether, triethyleneglycol methylvinyl ether, triethyleneglycol divinyl ether, trimethylol propane trivinyl ether, pentaerythritol triary ether, 1,4-cyclohexane dimethanol divinyl ether, and a mixture thereof.

The reactive diluent can control the cross-linking density of the composition to impart a good flexibility to the composition, and it may be used in an amount of 10 to 60 parts by weight in the composition.

In addition, the resin composition of the present invention comprises a silicone or fluorine group-containing compound ("Component C") as a functionalized additive. Component C has at least one silicon or fluorine group, and representative examples thereof include (i) a reactive monomer or oligomer having a silicone group, e.g., a silicone-containing vinyl derivative, silicone-containing (meth)acrylate, (meth)acryloxy-containing organosiloxane, or silicone polyacrylate; (ii) a reactive monomer or oligomer having a fluorine group, e.g., a fluoroalkyl-containing vinyl derivative, fluoroalkyl-containing (meth)acrylate, or fluorine polyacrylate; (iii) a silicone or fluorine containing resin, e.g., an organopolysiloxane, and a fluorinated polymer; and (iv) a silicone or fluorine containing surfactant or oil, e.g., a dimethyl silicone oil; and a mixture thereof.

The functionalized additive can provide a mold with a good releasing property, and it may be used in an amount of 5 to 200 parts by weight, in case of the ingredients (i) to (iii), and in an amount of 0.01 to 5 parts by weight, in case of the ingredient (iv), based on 100 parts of the total amount of Components A



and B used.

The photo-initiator ("Component D") used in the composition of the present invention may be a conventional free radical initiator or cationic initiator, or a mixture thereof. Representative examples of the free radical initiator include benzyl ketals, benzoin ethers, acetophenone derivatives, ketoxime ethers, benzophenone, benzo and thioxantone compounds, and mixtures thereof, and the cationic initiator may be onium salts, ferrocenium salts, diazonium salts, and mixtures thereof. In case a vinyl ether compound is used as Component B, it is preferred to employ a suitable mixture of the free radical initiator and the cationic initiator:

The photo-initiator may be preferably employed in an amount of 0.1 to 10 parts by weight based on 100 parts of the total amount of Components A, B and C used.

The resin composition of the present invention can be cured with an active energy ray to provide an organic mold having a good releasability, low swelling in solvent, good conformity to a substrate, and high mechanical strength. Further, the inventive resin composition can provide a mold having a large size in a simple process at a low production cost, and therefore, it can be utilized for the mass production of an organic mold.

The resulting organic mold can be beneficially used in forming an ultramicro- or submicro-pattern having a line width of below several ten nm. The formation of micropatterns using the inventive organic mold can be conducted by any replicating method known in the art, e.g., nano-imprint lithography, micro-contact printing (mCP), micro-molding in capillaries (MIMIC), micro-transfer molding (mTM), soft molding, and capillary force lithography (CFL).

FIG. 1 shows a schematic view of the procedure for preparing a mold using a resin composition according to the present invention. Specifically, as shown in the procedure (a), the inventive resin composition is coated or cast on a pattern face of a mastermold, and a support for the mold is covered thereon. The

resulting laminate is irradiated with an active energy ray such as a UV light to preliminarily cure the resin composition. The cured organic mold having a pattern face reverse to that of the mastermold is removed from the mastermold, and further UV-cured until the remaining reactive group in the resin is completely consumed, to enhance the hardness of the organic mold. The support for the mold may be preferably made of a polymer such as polyethylene terephthalate(PET), polycarbonate(PC), polyvinyl chloride(PVC), a soft or rigid elastomer, and others. The organic mold thus fabricated may be further adhered to or combined with a soft elastic or hard polymer backing having a desired shape and thickness, depending on the final use of the mold. The baking may be made of an epoxy resin, urethane elastomer, butadiene-based rubber, or a mixture thereof.

Alternatively, according to the procedure (b) of FIG. 1, a mastermold is firstly coated or cast with the resin composition of the present invention as in the (a) procedure and pseudo-cured with an UV light, and thereto an UV-curable or heat-curable resin composition as a backbone material is poured to a desired thickness in a vessel and completely cured. The cured organic mold is removed from the mastermold. The heat or UV curable backbone resin may be a conventional soft or rigid polymeric material selected from an epoxy resin, urethane elastomer, butadiene-based rubber, and a mixture thereof, depending on the final use of the mold.

The present invention will be described in further detail by the following Examples, which are, however, not intended to limit the scopes of the present invention.

### Preparations 1 to 5

Mold compositions having the composition shown in Table 1 were prepared.

Table 1

Component	Ingredient	Prep. 1	Prep. 2	Prep. 3	Prep. 4	Prep. 5
(A) Active energy curable urethane-based oligomer <sup>*1</sup>	Difunctional aliphatic urethane acrylate oligomer <sup>*5</sup>	31.0	37.5	31.0	31.0	36.0
	Difunctional cycloaliphatic urethane acrylate oligomer <sup>*6</sup>	25.0	-	25.0	25.0	20.0
	Trifunctional aliphatic urethane acrylate oligomer <sup>*7</sup>	12.5	25.0	12.5	12.5	27.5
	Trifunctional cycloaliphatic urethane acrylate oligomer <sup>*8</sup>	-	12.5	-	-	-
	Hexafunctional aliphatic urethane acrylate oligomer <sup>*9</sup>	6.5	-	6.5	6.5	11.5
(B) Reactive diluent <sup>*2</sup>	1,6-hexanediol diacrylate	12.5	12.5	12.5	12.5	-
	Neopentylglycol diacrylate	12.5	-	12.5	12.5	5.0
	Trimethylol propane (propylated) 3 triacrylate	-	12.5	-	-	-
	(3-acryloylpropyl)-tris(trimethylsiloxy)silane	12.5	12.5	-	-	-
(C) Functionalized additive <sup>*3</sup>	Silicone polyacrylate	12.5	-	-	-	-
	2,2,2-trifluoroethyl methacrylate	-	12.5	-	-	-
	Dimethyl silicone oil (100 centistokes)	-	-	1.0	-	1.0
	1-hydroxycyclohexyl phenyl ketone	2.5	2.5	2.5	2.5	2.5
(D) Free Radical Photoinitiator <sup>*4</sup>	Methyl benzoyl formate	1.5	1.5	1.5	1.5	1.5

Footnote:

\*1, \*2: part by weight

\*3: parts by weight based on the sum of Components A and B

\*4: parts by weight based on the sum of Components A, B and C

5 \*5: a urethane reaction product of hydroxypropyl acrylate and hexamethylene diisocyanate at an equivalent ratio of 2:1

\*6: a urethane reaction product of hydroxypropyl acrylate and 4,4-dicyclohexylmethane diisocyanate at an equivalent ratio of 2:1

10 \*7: a urethane reaction product of hydroxypropyl acrylate and a trifunctional triisocyanate having a structure of isocyanurate of hexamethylene diisocyanate, at an equivalent ratio of 3:1

\*8: a urethane reaction product of hydroxypropyl acrylate and a trifunctional triisocyanate having a structure of isocyanurate of isopropylidene diisocyanate, at an equivalent ratio of 3:1

15 \*9: a urethane reaction product of pentaerythritol propoxylate triacrylate and hexamethylene diisocyanate at an equivalent ratio of 2:1

### Example 1-1

20 An organic mold was fabricated by a replicating method as shown in the procedure (a) of FIG. 1.

Specifically, a silicon mastermold having the reverse pattern structure of a desired resin pattern was prepared. A resin composition according to Preparation 1 was coated on the pattern face of the mastermold to a thickness of 15  $\mu\text{m}$ . Then,  
25 a transparent adhesive polyethylene terephthalate sheet having a thickness of 188  $\mu\text{m}$  was laid on the coated surface, and the resulting laminate was irradiated with a UV light of 5  $\text{mJ}/\text{cm}^2$  for 15 seconds to cure the resin composition.

The cured organic mold was lifted off from the mastermold to obtain an organic mold having the desired pattern combined with a polyethylene  
30 terephthalate (PET) support. Then, UV (5  $\text{mJ}/\text{cm}^2$ ) was additionally irradiated

for 2 hours to completely cure the organic mold.

### Example 1-2

5 A replicated organic mold was fabricated as shown in the procedure (b) of FIG. 1.

Specifically, the pattern surface of the silicon mastermold used in Example 1-1 is firstly coated with the resin composition of Preparation 1 to a thickness of 15  $\mu\text{m}$  and pre-cured for 3 minutes with a UV light of 5  $\text{mJ}/\text{cm}^2$ .

10 The laminate was placed in a vessel, and a UV curable, acrylated butadiene resin composition was poured and cured to form a backbone layer to a thickness of 2 mm.

The cured product was removed from the vessel and an organic mold having the desired pattern with an acrylated butadiene backbone was lifted off  
15 from the mastermold, and then additionally irradiated with UV(5  $\text{mJ}/\text{cm}^2$ ) for 2 hours to completely cure the organic mold.

FIG. 2a is an electronic microscopic photograph of an organic mold fabricated as in Example 1-1 which has an equal line width and space of 80 nm.

20 Further, FIG. 3a is an electronic microscopic photograph of an organic mold having a cylindrical pattern, in which cylindrical cavities having a 100 nm diameter and a 450 nm height were aligned in series, fabricated as in Example 1-1.

### Example 2-1

25

The organic mold shown in FIG. 2a was employed to form a pattern by a soft molding method. Specifically, a polystyrene resin solution was coated on a silicon wafer substrate and, on the coated resin layer, the mold was placed and slightly pressed to transfer the desired pattern on the polystyrene resin layer. FIG.

30 2b shows an electronic microscopic photograph of the pattern thus formed.

### Example 2-2

The organic mold shown in FIG. 3a was employed to form a pattern by  
5 UV flash replication method. Specifically, a UV-curable, acrylated epoxy resin  
solution was coated on a PET substrate, the prepared mold was placed on the  
coated epoxy resin layer, and then the resulting laminate was irradiated with UV  
to cure the epoxy resin composition. FIG. 3b is an electronic microscopic  
photograph of the pattern thus formed.

10 FIGs. 2b and 3b illustrate that the mold fabricated from the inventive  
resin composition can provide submicropatterns without the problem of adhesion  
or generating defects.

### 15 Examples 3 and 4

The procedure of Example 1-1 was repeated except for using the mold  
composition of Preparation 2 and 3 instead of the composition of Preparation 1,  
respectively, to fabricate organic molds, which were then used in forming patterns  
20 according to Example 2-1.

The organic molds thus fabricated and the patterns obtained therefrom  
have the same characteristics observed in FIGs. 2a and 3a, and 2b and 3b,  
respectively.

### 25 Comparative Example 1

The procedure of Example 1-1 was repeated except that the resin  
composition of Preparation 4 which contains no functionalized additive, and a  
silicon mastermold having a pyramid pattern, in which pyramid shaped cavities  
30 having a bottom area of  $180\ \mu\text{m} \times 180\ \mu\text{m}$  and a height of  $70\ \mu\text{m}$  were aligned in

series; was used to fabricate an organic mold.

FIG. 4a is an optical microscopic photograph of the pyramid patterns-forming mastermold, and FIG. 4b is an optical microscopic photograph of an organic mold fabricated using the mastermold. By comparing FIG. 4a and FIG. 4b, it can be seen that the organic mold thus fabricated has several defects.

### Comparative Example 2

The procedure of Example 1-1 was repeated except that the organic mold composition of Preparation 5 comprising a smaller amount of a reactive diluent was used to fabricate an organic mold. The organic mold broke easily during the patterning step due to its high brittleness.

While the invention has been described with respect to the above specific examples, it should be recognized that various modifications and changes may be made to the invention by those skilled in the art which also fall within the scope of the invention as defined by the appended claims.

What is claimed is:

1. A resin composition for a mold used in forming micropatterns, which comprises:
  - 5 (A) 40 to 90 parts by weight of an active energy curable urethane-based oligomer having a reactive group selected from the group consisting of (meth)acrylate, vinyl ether, arylether, and a combination thereof;
  - (B) 10 to 60 parts by weight of a monomer reactive with the urethane-based oligomer, having a reactive group selected from the group  
10 consisting of (meth)acrylate, vinyl ether, arylether, and a combination thereof;
  - (C) 0.01 to 200 parts by weight of a silicone or fluorine containing compound, based on 100 parts of the sum of the components (A) and  
15 (B); and
  - (D) 0.1 to 10 parts by weight of a photoinitiator, based on 100 parts of the sum of the components (A), (B) and (C).
2. The composition according to claim 1, wherein the active energy curable  
20 urethane-based oligomer is selected from the group consisting of linear aliphatic, cycloaliphatic and aromatic urethane-based oligomers having at least two reactive groups, and a mixture thereof.
3. The composition according to claim 1, which further comprises at least one  
25 reactive oligomer selected from the group consisting of a (meth)acrylated polyester, (meth)acrylated polyether, (meth)acrylated epoxy, (meth)acrylated polycarbonate, (meth)acrylated polybutadiene, and a mixture thereof, as a substituent of Component A.
- 30 4. The composition according to claim 1, wherein the (meth)acrylate used as



Component B is selected from the group consisting of isobornyl acrylate, 1,6-hexanediol acrylate, triethyleneglycol di(meth)acrylate, trimethylol propane triacrylate, tetraethyleneglycol di(meth)acrylate, 1,3-butanediol diacrylate, 1,4-butanediol diacrylate, diethyleneglycol diacrylate, neopentylglycol diacrylate, neopentylglycol di(meth)acrylate, polyethyleneglycol di(meth)acrylate, pentaerythritol triacrylate, dipentaerythritol (hydroxy) pentaacrylate, alkoxylated tetraacrylate, octadecyl acrylate, isodecyl acrylate, lauryl acrylate, stearyl acrylate, behenyl acrylate, styrenic monomer, and a mixture thereof.

5. The composition according to claim 1, wherein the vinyl ether used as Component B is selected from the group consisting of cyclohexyl vinyl ether, 2-ethylhexyl vinyl ether, dodecyl vinyl ether, 1,4-butanediol divinyl ether, 1,4-hexanediol divinyl ether, diethylene glycol divinyl ether, ethyleneglycol butyl vinyl ether, ethyleneglycol divinyl ether, triethyleneglycol methylvinyl ether, triethyleneglycol divinyl ether, trimethylol propane trivinyl ether, 1,4-cyclohexane dimethanol divinyl ether, and a mixture thereof.

6. The composition according to claim 1, wherein the aryl ether used as Component B is selected from the group consisting of aryl propyl ether, aryl butyl ether, pentaerythritol triary ether, and a mixture thereof.

7. The composition according to claim 1, wherein the silicone or fluorine-containing compound is at least one selected from:

- (i) a silicone-containing reactive compound selected from the group consisting of a silicone-containing vinyl derivative, silicone-containing (meth)acrylate, (meth)acryloxy-containing organosiloxane, silicone polyacrylate, and a mixture thereof;
- (ii) a fluorine-containing reactive compound selected from the group consisting of a fluoroalkyl-containing vinyl derivative, fluoroalkyl-containing (meth)acrylate, fluorine polyacrylate, and a mixture

thereof;

- (iii) a silicone or fluorine containing resin, or a mixture thereof; and
- (iv) a silicone or fluorine containing surfactant or oil, or a mixture thereof.

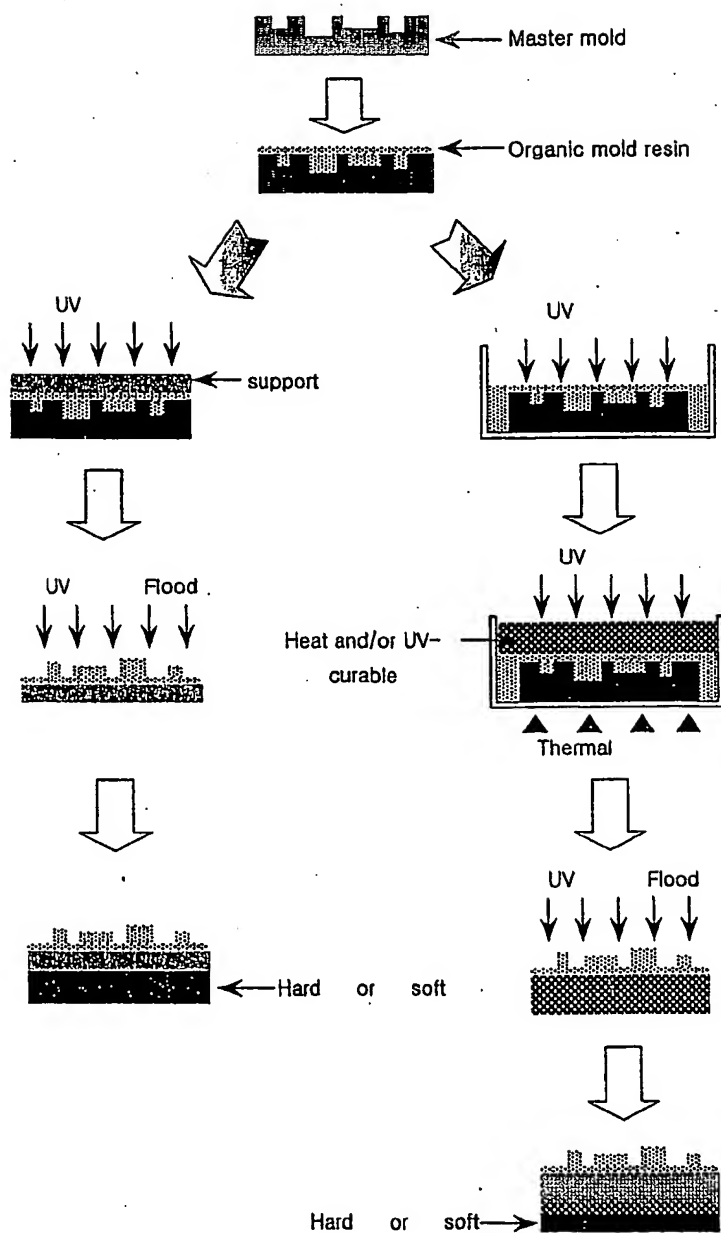
5 8. The composition according to claim 1, wherein the photoinitiator is at least one of a free radical initiator selected from the group consisting of benzyl ketals, benzoin ethers, acetophenone derivatives, ketoxime ethers, benzophenone, benzo and thioxantone compounds, and mixtures thereof, and a cationic initiator selected from the group consisting of onium salts, ferrocenium salts, diazonium  
10 salts, and mixtures thereof.

9. A method for fabricating an organic mold, which comprises coating or casting the resin composition recited in claim 1 on a pattern face of a mastermold, placing a support on the resin layer, irradiating the resulting laminate with an active  
15 energy ray to preliminarily cure the resin composition, lifting off the organic mold having a reverse pattern face to that of the mastermold and integrally formed with the support from the mastermold, and completely curing the organic mold.

10. The method according to claim 9, which further comprises adhering a soft or  
20 rigid backing having a curved or flat face to the bottom face of the organic mold.

11. A method for fabricating an organic mold, which comprises coating or casting a resin composition recited in claim 1 on a pattern face of a mastermold, irradiating the resin layer with an active energy ray to preliminarily cure it, pouring a UV- or heat-curable resin composition onto the cured resin layer as a  
25 backbone, heating or irradiating the resultant to completely cure the resin and the backbone layers, lifting off the organic mold having a reverse pattern face to that of the mastermold and integrally formed with the backbone layer from the mastermold, and completely curing the organic mold.

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FIG. 1





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FIG. 2a

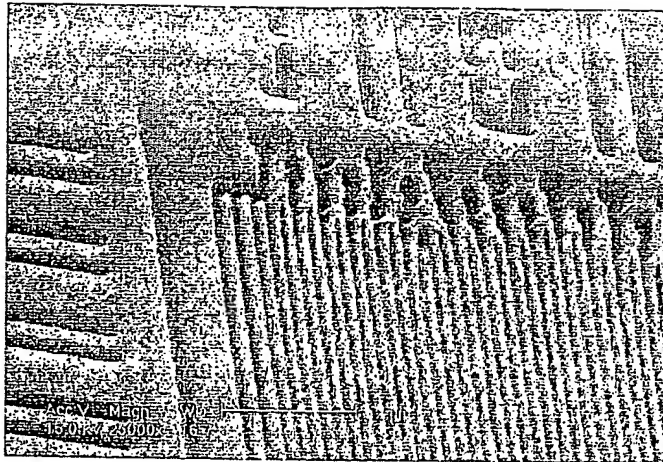
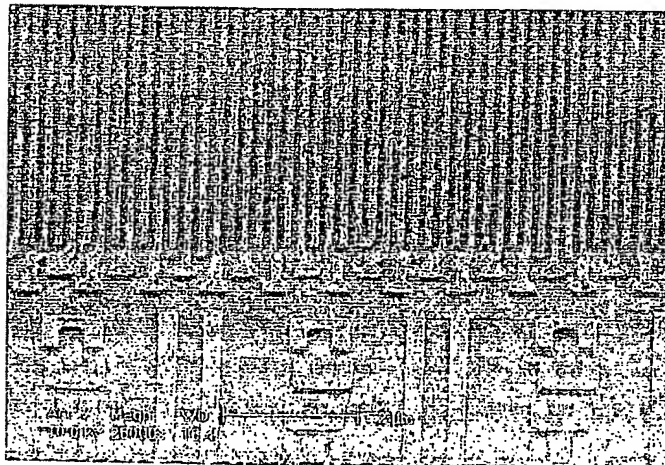


FIG. 2b





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FIG. 3a

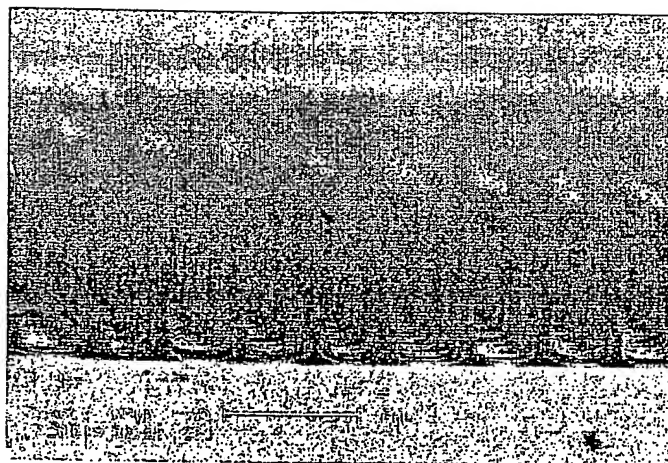
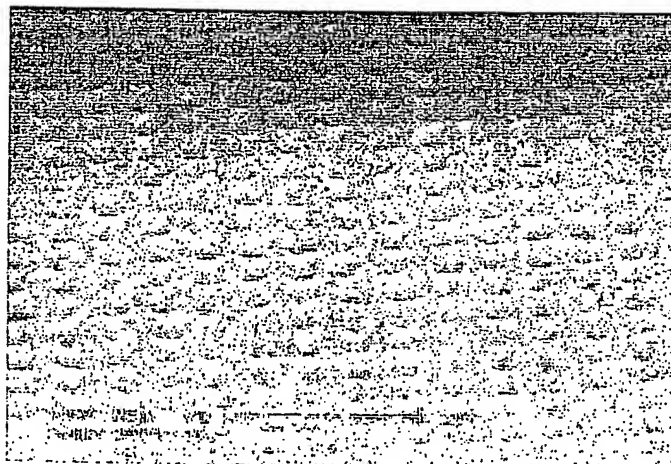


FIG. 3b





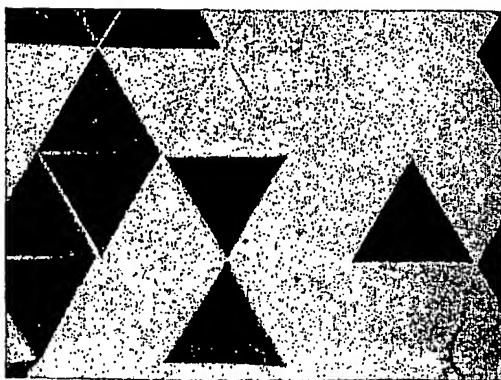


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FIG. 4a



FIG. 4b





## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/KR2004/000860

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC7 G03F 7/027

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

IPC7 G03F 7/027, G02B 3/00, B05D 5/06, C08G 18/42, B32B27/00, C08F 2/44, B41N 1/08

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
KR, JP : classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
NPS, DELPHION, ESPACENET, USFTO, JPO

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 882 998 A1 (DAI NIPPON PRINTING CO., LTD) 09. Dec. 1998 see the whole document	1 - 2, 7
Y		3 - 6, 8 - 11
Y	EP 421 027 A1 (SEKISUI KAGAKU KOGYO) 10. Apr. 1991 see the whole document	3 - 6
Y	JP 1995-318706 A (DAI NIPPON PRINTING CO., LTD) 08. Dec. 1995 see the whole document	8 - 11
Y	JP 1994-201903 A (MITSUBISHI CO., LTD) 22. Jul. 1994 see the whole document	8 - 11
Y	JP 1994-118205 A (MITSUBISHI CO., LTD) 28. Apr. 1994 see the whole document	8 - 11

☐ Further documents are listed in the continuation of Box C.

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Date of the actual completion of the international search  
22 JULY 2004 (22.07.2004)

Date of mailing of the international search report  
29 JULY 2004 (29.07.2004)

Name and mailing address of the ISA/KR



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# INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/KR2004/000860

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 882 998 A1	09. Dec. 1998	US 6071443 A CA 2243114 AA WO 9823978 A1 DE 69714972 C0	06. Jun. 2000 04. Jun. 1998 04. Jun. 1998 02. Oct. 2002
EP 421 027 A1	10. Apr. 1991	none	
JP 1995-318706 A	08. Dec. 1995	none	
JP 1994-201903 A	22. Jul. 1994	none	
JP 1994-118205 A	28. Apr. 1994	none	

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